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## Abstract

The present invention relates to novel compounds of formula (I):

wherein

---- represents a single or a double bond;

R is a radical selected from:

$$i)^{(R_1)p}$$
  $ii)$   $(R_1)p$   $iii)$   $(R_1)p$  and  $iv)$   $(R_1)p$ 

in which R<sub>1</sub> is halogen, cyano, C<sub>1-4</sub> alkyl, C<sub>1-4</sub> alkoxy, trifluoromethyl or trifluoromethoxy and p is zero or an integer from 1 to 3;

R<sub>2</sub> is hydrogen or C<sub>1-4</sub> alkyl;

R<sub>3</sub> is hydrogen, hydroxy or C<sub>1-4</sub> alkyl;

R<sub>4</sub> is hydrogen or R<sub>4</sub> together with R<sub>3</sub> represents =0 or =CH2;

R5 is phenyl, naphthyl, a 9 to 10 membered fused bicyclic heterocyclic group or a 5 or 6 membered heteroaryl group, wherein said groups are optionally substituted by 1 to 3 groups independently selected from trifluoromethyl, C<sub>1-4</sub> alkyl, hydroxy, cyano, C<sub>1-4</sub> alkoxy, trifluoromethoxy, halogen or S(O)qC<sub>1-4</sub> alkyl;

 $R_{6}$  and  $R_{7}$  independently are hydrogen, cyano,  $C_{1\text{--}4}$  alkyl;

 $R_8$  is (CH<sub>2</sub>)rR<sub>10</sub>;

Rg is hydrogen, halogen,  $C_{3-7}$  cycloalkyl, hydroxy, nitro, cyano or  $C_{1-4}$  alkyl optionally substituted by one or two groups selected from halogen, cyano, hydroxy or  $C_{1-4}$  alkoxy;

R<sub>10</sub> is hydrogen or C<sub>3-7</sub> cycloalkyl;

n is 1 or 2;

q is 0, 1 or 2;

r is 0 or an integer from 1 to 4;

PB60564U\$W

or a pharmaceutically acceptable salt or a solvate thereof, process for their preparation and their use in the treatment of conditions mediated by tackykinins and/or by selective inhibition of the serotonin reuptake transporter protein.